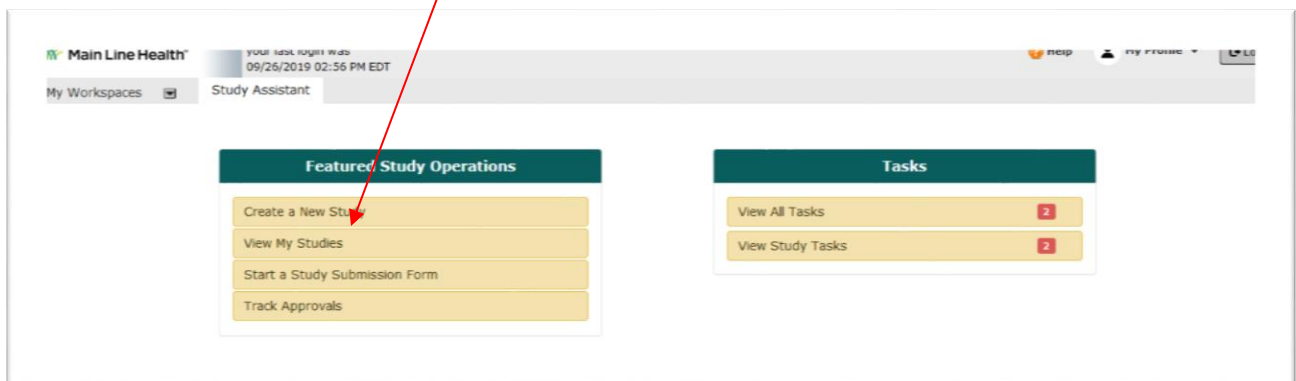


# Stamped Documents

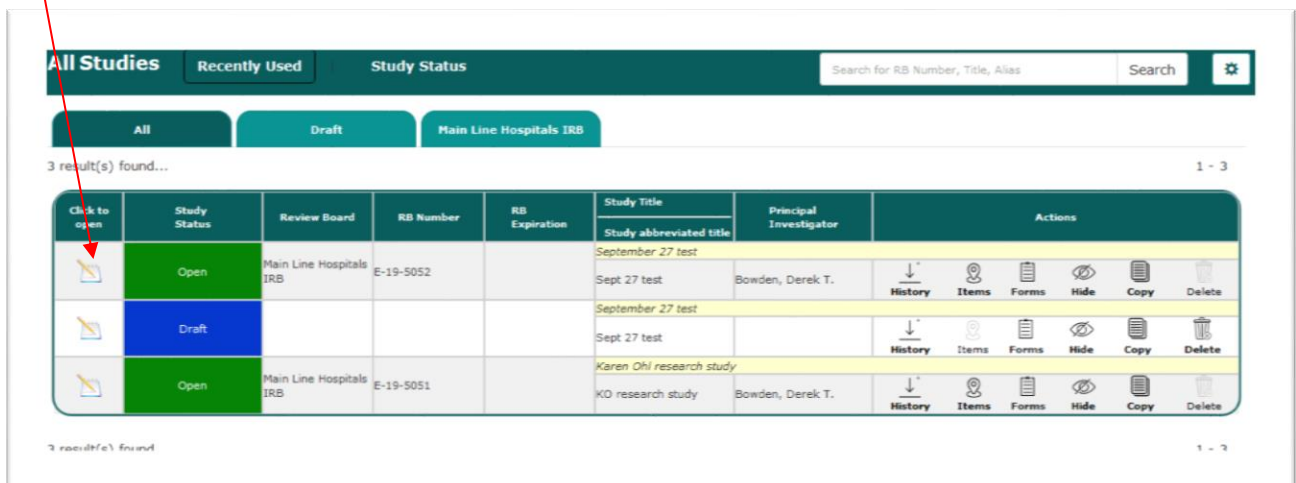


## Accessing Study Documents after MLH IRB Approval

Once you receive notification that the MLH IRB has Approved your submission, log into imedris and find your approved submission: click "View My Studies"



Click to open the approved study.



# MAIN LINE HEALTH OFFICE OF RESEARCH PROTECTIONS

You will be taken to the Study dashboard. You can access the study application, generate new forms to submit for this study, and access the stamped consent and other study documents.

The screenshot shows the 'Study Management' dashboard for a study with IRB Number E-19-5052 and Study Title 'September 27 test'. The 'Protocol Items' section includes links for 'Study Application', 'Informed Consent', 'Other Study Documents', 'Adverse Event Reporting Form', 'Amendment/General Reporting Form', 'Initial Submission Packet', and 'Continuing Review Submission Form'. A red arrow points from the 'Informed Consent' link to the second screenshot. The 'Outstanding Submission(s)' table lists two items:

Track Location	Ref Number	Request Type	Process Submission
	000056	Click on the hyperlink to edit/view the submission. Amendment/General Reporting Form	Send Submission
Routing In Process	000055	Click on the hyperlink to edit/view the submission. Continuing Review Submission Form	Retract Submission

Clicking into Informed Consent will allow you to download the IRB approved stamped consent.

The screenshot shows the 'Informed Consent' management interface. It includes search filters for 'Search Level', 'Select Category', 'Version #', and 'Approval Date'. A table of consent records is displayed below:

View History	Edit/View	Title/Category	Version	Language	Un-Approved Consent	Approved Consent	Consent Outcome	Approval Date	Expiration Date	Checked Out By	Create a Revised Document
		Consent	1.0 09/27/2019	English			Approved	09/27/2019	09/26/2020		Add Revision

**Questions?** Contact the Main Line Health Office of Research Protections at 610.225.6222.